



PATENT ABSTRACTS OF JAPAN

(11) Publication number: **10330369 A**(43) Date of publication of application: **15.12.98**

(51) Int. Cl.

C07D263/58
A61K 31/415
A61K 31/42
A61K 31/425
A61K 31/435
A61K 31/44
C07D235/30
C07D277/82
C07D401/12
C07D413/12
C07D417/12
C07D471/04
C07D473/00
C07D473/40
C07D498/04
C07D513/04

(21) Application number: **09225679**(22) Date of filing: **06.08.97**(30) Priority: **04.04.97 JP 09102481**

(71) Applicant:

SUMITOMO PHARMACEUT CO
LTD SUMITOMO CHEM CO LTD

(72) Inventor:

TOKUNAGA TERUHISA
FUJITA ICHIKI
NAGATA TATSU
OCHI HIROSHI
WATANABE KOSEI
NAKATANI TOMOSUKE

(54) HETEROCYCLIC COMPOUND

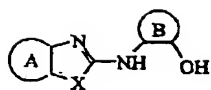
(57) Abstract:

PROBLEM TO BE SOLVED: To obtain the subject composition, capable of suppressing immune response of a type 2 helper T cell (Th2) and useful as a therapeutic agent for allergic diseases, systemic lupus erythematosus, AIDS, etc., by using a specific heterocyclic compound as an active ingredient.

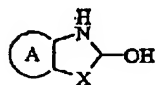
SOLUTION: This pharmaceutical composition contains a heterocyclic compound represented by formula I [X is O, NH or S; the ring A is a (substituted) benzene ring or a 6-membered ring (substituted) heterocyclic ring containing one or two N atoms; the ring B is a(n)

(electron attractive group-substituted) benzene ring or a(n) (electron attractive group-substituted)pyridine ring] or its pharmaceutically acceptable salt [e.g. 2-(6-nitro-2-benzoxazolyl)amino-5-nitrophenol] as an active ingredient. The compound is preferably obtained by reacting a compound, prepared from, e.g. an amine derivative as a starting raw material and represented by formula II or III with an amino-alcohol derivative represented by formula IV in an inert organic solvent at a temperature within the range of about ambient temperature to about the boiling point of the solvent while suppressing the reaction.

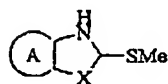
COPYRIGHT: (C)1998,JPO



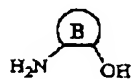
I



II



III



IV